

Neuro-ophthalmology review

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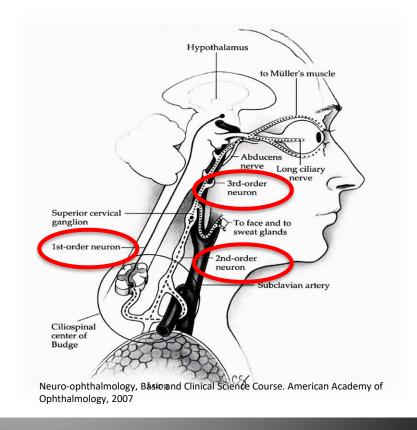
Neuro-ophthalmology deals with visual problems caused by disorders of the brain or the optic nerve connection

Part 1:

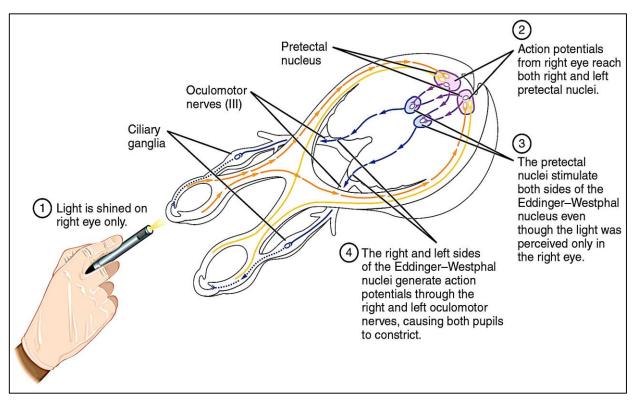


- Anatomy and physiology:
 - The pupil size of is controlled by a balance between parasympathetic innervation to the sphincter muscles and sympathetic innervation of the dilator muscles of the iris.
 - Pupil constrict to light and near stimuli.

- Sympathetic (adrenergic) pathway:
 - Pupillary dilation is mediated through sympathetic (adrenergic) pathway that originate in the hypothalamus



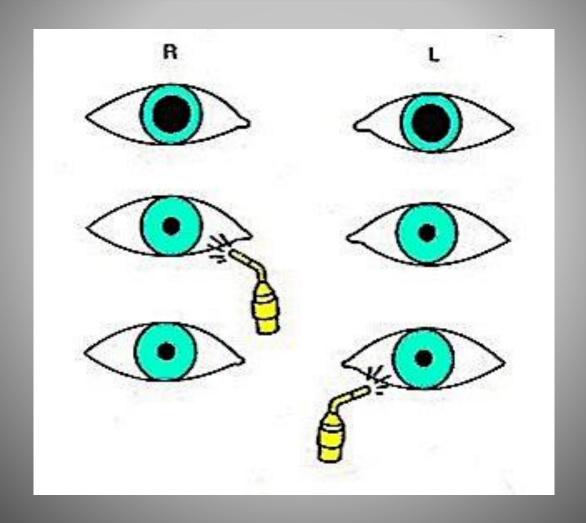
parasympathetic (cholinergic) pathway:

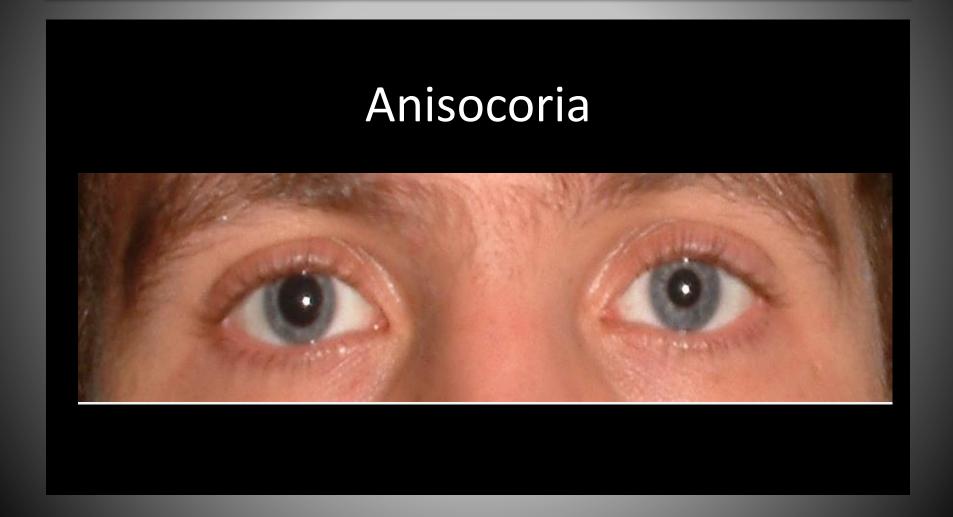


http://oerpub.github.io/epubjs-demo-book/resources/1509_Pupillary_Reflex_Pathways.jpg

- Examination of the pupil :
 - Best conducted in dim light room using a bright light
 - The patient should be relaxed and fixing on a distant object
 - The size, shape, and position of each pupil should be noted in light and dark conditions
 - Check light reflex looking for a relative afferent pupillary defect (RAPD)

Pupillary light reflex





Which pupil is abnormal?



When the small pupil does not dilate as well as the large pupil in dim light, then the small pupil is abnormal.

 When the larger pupil does not constrict as well as the small pupil in response to a light stimulus, then the large pupil is abnormal.

The large pupil is abnormal

- Previous ocular surgery
- Ocular trauma
- Use of medication like cycloplegics e.g. atropine, cyclopentolate
- Third nerve palsy
- Tonic pupil (Adie's pupil)

Tonic pupil (Adie's pupil)

- Sluggish, segmental pupillary responses to light
- better response to near followed by slow redilation.
- Young female
- Unilateral (80%)
- Instillation of weak cholinergic agents (0. 1% pilocarpine) will cause constriction of the tonic pupil (denervation hypersensitivity)
- Benign condition

Tonic pupil (Adie's pupil)



Holmes-Adie syndrome:

 includes tonic pupil, diminished deep tendon reflexes and orthostatic hypotension.

The small pupil is abnormal:

- Previous ocular surgery
- Ocular trauma or inflammation
- Use of medication e.g. pilocarpine
- Horner syndrome



The small pupil is abnormal:

Horner syndrome:



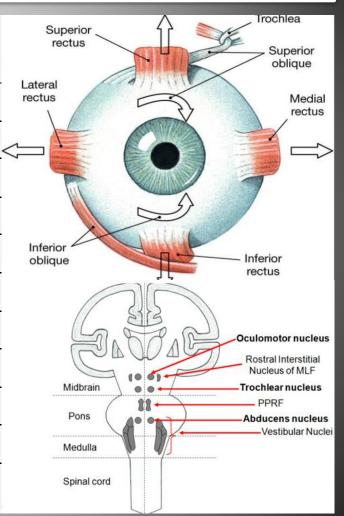
- Small pupil (miosis), ptosis and anhydrosis
- Caused by a lesion anywhere along the sympathetic pathway
- Carotid dissection, carotid aneurysm and tumor can be associated with this syndrome

Part 2:

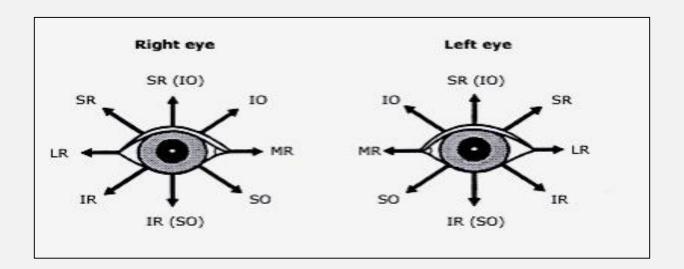
Neuromotility disorders

Anatomy and physiology:

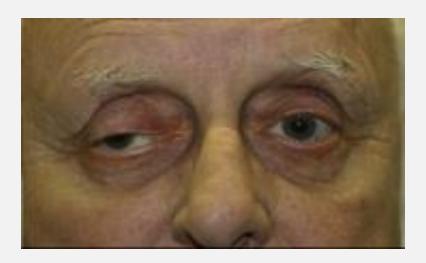
Innervation@f@xtraocular@nuscles@	Primary@ction@
Cranial nerve III 2	
Superior@ectus@	Elevation@maximal@n@ateral@aze)@
Inferior@ectus2	Depression[[maximal@nfateralgaze]]
Medial@ectus@	Adduction
Inferior@blique@	Excyclotorsion
Cranial nerve IV2	
Superior@blique2	Incyclotorsion 2
Cranial nerve VI	
Lateral@ectus@	Abduction
2	



Anatomy and physiology:



 65 yrs old presented to ER complaining of double vision



Third cranial nerve (oculomotor)palsy :



Neurosurg Clin N Am 23 (2012) 607-619

Third cranial nerve (oculomotor)

- Begins as a nucleus in the midbrain that consists of several subnuclei.
- innervate the individual extraocular muscles, the eyelids, and the pupils

Third cranial nerve (oculomotor)palsy:

Check for pupil involvement

Absence of pupillary involvement suggests a benign process that can be observed over a couple of weeks. A fixed, dilated pupil requires extensive neurologic evaluation.

Third cranial nerve (oculomotor)palsy:

Check for pupil involvement





Third cranial nerve (oculomotor)palsy :

- Etiology:
- Micro-vascular ischemia (DM and HTN)
- Intracranial aneurysm (posterior communicating artery)
- Trauma
- Brain tumor

- Fourth cranial nerve (trochlear) palsy:
 - Vertical diplopia
 - Head tilt to the opposite shoulder

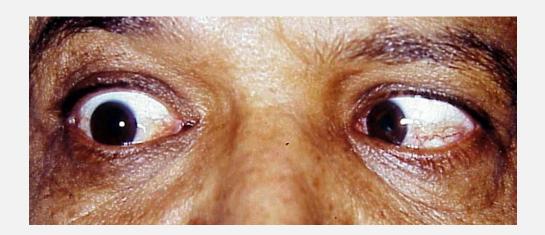


Fourth cranial nerve (trochlear) palsy:

- Etiology:
 - trauma
 - idiopathic
 - -congenital

Which muscle is affected?





Sixth cranial nerve (abducens)palsy :

- Horizontal diplopia (worse at distance)
- Esotropia
- Face turn in the direction of the paralyzed muscle
- Limited Abduction on the side of the lesion

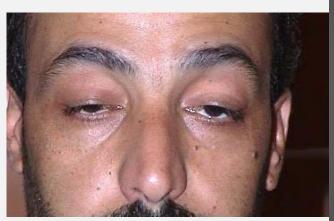
- Sixth cranial nerve (abducens)palsy:
 - causes:
 - -intracranial tumors
 - trauma
 - microvascular diseases
 - increased intracranial pressure

Part 3:

Neuromuscular disorder

Ocular myasthenia gravis

- Chronic autoimmune disease affecting the neuromuscular junction in skeletal muscles.
- Ptosis
- Diplopia
- Fatigability and variability of clinical findings are characteristic
- The pupil is not affected



Ocular myasthenia gravis

- Check for systemic weakness, difficulty in swallowing or breathing.
- Assess orbicularis strength
- Blood test for acetylcholine receptor antibodies

Ocular myasthenia gravis:

Tensilon test: inhibits acetylcholinesterase and can transiently reverse signs of weakness due to OMG, such as ptosis and extraocular muscle paresis.

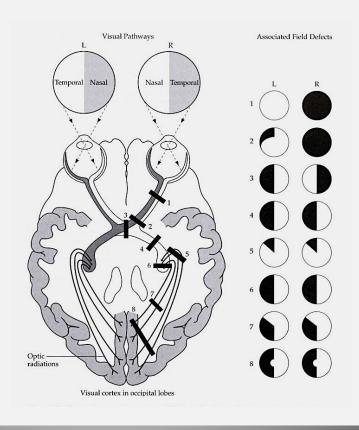
Part 4:

Visual pathway disorders

Visual pathway disorders:

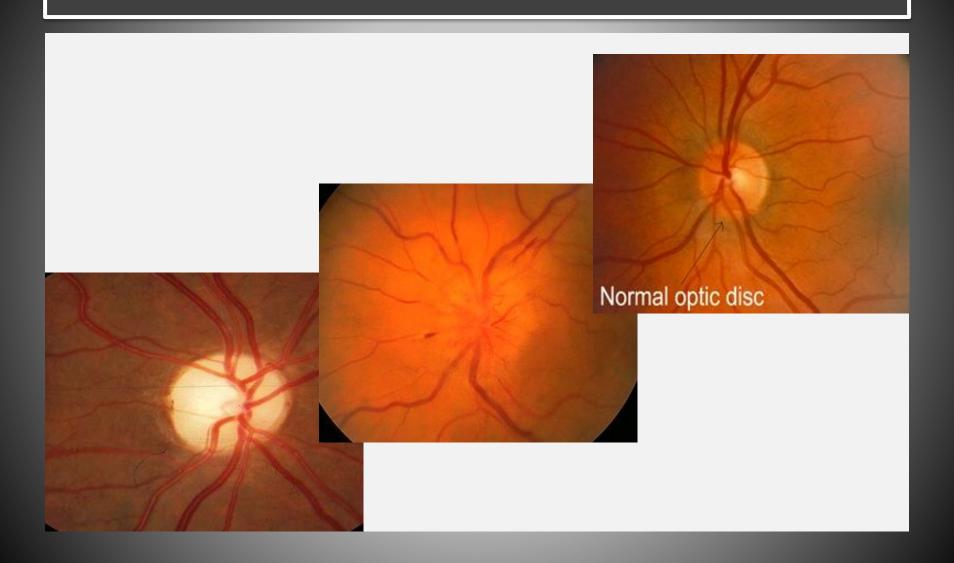
Lesions anywhere in the visual pathway will produce

visual field defect



Optic nerve disease:

- Usually unilateral
 - Afferent pupillary defect
 - Central visual loss
 - -Loss of color vision
 - Optic disc edema
 - Optic atrophy



Typical optic neuritis :

- Inflammatory demyelinating condition associated with MS
- Most common type in young adults
- Good recovery
- IV steroids my speed up the recovery process but does not influence the final outcome.

Ischemic optic neuropathy (ION):

- Non-arteritic ION:
- Patients often have DM,HTN and other vascular risk factor.
- Most common cause in older patients
- Altitudinal visual field loss

Visual pathway disorders

Ischemic optic neuropathy (ION):

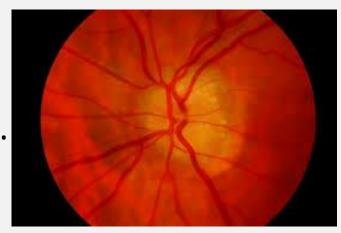
- Arteritic ION:
- >55yrs old
- Associated with giant cell arteritis (GCA)
- Check for jaw claudication, proximal myalgia and arthralgia, scalp tenderness, headache
- Elevated erythrocyte sedimentation rate (ESR) and Creactive protein (CRP)

Arteritic ION:

- Temporal artery biopsy is the gold standard for diagnosis
- Treatment: Systemic steroids is given immediately if GCA is suspected
- Binocular involvement occurs in third of cases, often within the first day.

Congenital disc elevation: <1%

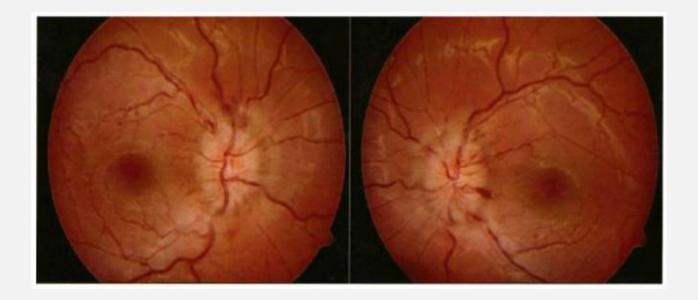
- -Optic disc margins blurred and the cup is absent but no edema or hrg can be observed.
- -May be associated with hyperopia or drusen



- Other causes of optic neuropathy:
 - Infection e.g viruses, TB, cryptococcus and syphilis
 - Systemic connective tissue disease e.g SLE
 - genetics: Leber's optic neuropathy (through a mitochondrial DNA mutation)
 - Toxic and nutritional deficiencies
 - Trauma

Papilledema

 Bilateral swelling of the optic discs secondary to increased intracranial pressure



Papilledema

- Hyperemia of the disc
- -Tortuosity of the veins and capillaries
- Blurring and elevation of disc margins
- Peripapillary flameshaped haemorrhages



Papilledema

Look for spontaneous venous pulsations

Visual pathway disorders

Papilledema

Causes:

- Intracranial mass
- Sever systemic hypertension
- Idiopathic intracranial hypertension (pseudotumor cerebri)

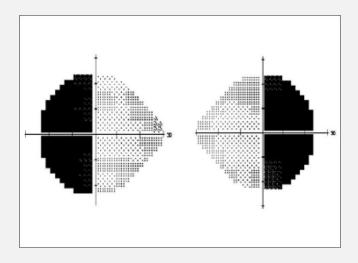
MCQ

a patient presented with this visual field defect.

Which one of the following diagnosis is the most

Likely?

- a. Optic neuritis
- b. tilted discs
- c. pituitary tumor
- d. 6th nerve palsy



Thank you